PHICESOEUGY.

JUL 1 9 1999

07/505977 8/13/90
Group Art Unit | Examiner

Paper No.

Assistant Commissioner for Patents Washington, DC 20221

hereby request eccess undent? CFR 1.14(a)(3)(iv) to the demilied AEANDONED application, which is: (CHECK C	te application file record of the acove. NE:
(A) referred to in United States Patent Number 50	323065 <u>ceiuma</u>
(B) referred to in an application that is open to opplication No filed	respection as set forth in 37 CFR !!!
(C), an application that calins the benefit of the filling inspection, i.e., Application No.	
(D) an application in which the applicant has filed a application to the public.	n authorization to lay open the como:
Please direct any correspondence concerning this rec	uest to the following address:
	5/6/99
Signature	Date
Billy Mullenap	FOR FTO USE SALY
Typed or unnied name	Approved by (Inc.
•	Unit:



United States Patent [19]

Cook et al.

[11] Patent Number:

5,623,065

[45] Date of Patent:

Apr. 22, 1997

[54] GAPPED 2' MODIFIED OLIGONUCLEOTIDES

[75] Inventors: Phillip D. Cook, Vista; Brett P. Monia, Carlsbad, both of Calif.

[73] Assignee: Isis Pharmaceuticals, Inc., Carlsbad,

Calif.

[21] Appl. No.: 244,993

[22] PCT Filed: Dec. 23, 1992

[86] PCT No.: PCT/US92/11339

§ 371 Date: **Jun. 21, 1994** § 102(e) Date: **Jun. 21, 1994**

[87] PCT Pub. No.: WO93/11339

PCT Pub. Date: Jul. 8, 1993

Related U.S. Application Data

[63]	Continuation-in-part of Ser. No. 814,961, Dcc. 24, 1991	,
	abandoned, and Ser. No. 566,977, Aug. 13, 1990, aban-	-
	doned.	_

[51]	Int. Cl.6	 C07H 21/00;	C07H	21/02;
			C07H	21/04

935/6; 935/9; 935/10=

6, 10

[56] References Cited

U.S. PATENT DOCUMENTS

4,867,187	9/1989	Duck 435/6
4,908,307	3/1990	Rodland et al 435/6
5,013,830	5/1991	Ohtsuka et al 536/25.1
5,034,506	7/1991	Summerton et al 528/391
5,134,066	7/1992	Rogers et al 435/91.3
5,149,797	9/1992	Pederson et al 536/23.1
5,220,007	6/1993	Pederson et al 536/23.1
5,256,775	10/1993	Froehler 536/25.6
5,366,878	11/1994	Pederson et al 435/91.3
5,403,711	4/1995	Walder et al 435/6
5,466,786	11/1995	Buhr et al 536/26.26

FOREIGN PATENT DOCUMENTS

FOREIGN PATENT DUCUMENTS				
11/1990	Canada .			
8/1987	European Pat. Off			
3/1989	European Pat. Off. ,			
4/1989	European Pat. Off			
11/1990	European Pat. Off			
9/1990	Germany .			
10/1992	Germany .			
of 1991	Japan .			
6/1989	WIPO.			
6/1990	WIPO .			
10/1990	WIPO.			
4/1991	WIPO .			
8/1991	WIPO .			
2/1994	WIPO.			
9/1994	WIPO .			
	11/1990 8/1987 3/1989 4/1989 11/1990 9/1990 10/1992 of 1991 6/1989 6/1990 10/1990 4/1991 8/1991 2/1994			

OTHER PUBLICATIONS

Block et al. 1988 Gene 72, 349-360.

Cormier et al. 1988 Nuc. Acids Res. 16(10), 4583–4594. Uhlmann et al. 1990 Chemical Reviews 90(4), 544–584.

Ikehara et al. 1977 Nuc. Acids Res. 4(12): 4249-4260.

Berkowitz et al. 1973 J. Medicinal Chemistry, 16(2):

<u>813-814.</u>

Kawasaki et al. 1991 (Jan.) "Synthesis and Biophysical Studies of 2'-dRIBO-F Modified Oligonucleotides", Conference on Nucleic Acid Therapeutics, Clearwater, FL.

Agrawal, S. et al., "Oligodeoxynucleoside Phosphoramidates and Phosphorothioates as Inhibitors of Human Immunodeficiency Virus" *Proc. Natl. Acad. Sci. USA* 1988 85, 7079–7083.

Augustyns, et. al., "Influence of the Incorporation of (S)-9-(3,4-dihydroxy-butyl)Adenine on the Enzymatic Stability and Base-Pairing Properties of Oligodeoxynucleotides" *Nucleic Acids Research* 1991, 19, 2587-2593.

Beaton, et. al., Chapter 5, Synthesis of oligonucleotide phosphorodithioates, p. 109, Oligonucleotides and Analogs, A Practical Approach, Eckstein, F., Ed.; The Practical Approach Series, IRL Press, New York, 1991, pp. 109–135. Borthwick, et al., "Synthesis of Chiral Carbocylic Nucleosides": Tetrahedron 1992, 48, 571–623.

Brill et al., "Synthesis of Deoxydinucleoside Phosphorodithioates", J. Am. Chem. Soc. 1991 113, 3972-3980.
Cohen in Oligonucleotides: Antisense Inhibitors of Gene

Expression, CRC Press, Inc., Boca Raton, Fl (1989), pp. 1-255.

Dagle et al., "Physical properties of oligonucleotides containing phosphoramidate-modified internucleoside linkages", Nucleic Acids Research 1991 19, 1805–1810.

Dagle et al., "Targeted degradation of mRNA in Xenopus oocytes and embryos directed by modified oligonucleotides: studies of An2 and cyclin in embryogenesis", Nucleic Acids Research 1990 18, 4751–4757.

Dagle et al., "Pathways of Degradation and Mechanism of Action of Antisense Oligonucleotides in Xenopus laevis Embryos", Antisense Research and Development 1991 1.

Debart et al., "Intermolecular Radical C—C Bond Formation: Synthesis of a Novel Dinucleoside Linker for Non-anionic Antisense Oligonucleosides", *Tetra. Ltrs.* 1992 33, 2645-2648.

(List continued on next page.)

Primary Examiner—Christopher S. F. Low Attorney, Agent, or Firm—Woodcock Washburn Kurtz Mackiewicz & Norris

[57] ABSTRACT

Oligonucleotides and other macromolecules are provided that have increased nuclease resistance, substituent groups for increasing binding affinity to complementary strand, and subsequences of 2'-deoxy-erythro-pentofuranosyl nucleotides that activate RNase H enzyme. Such oligonucleotides and macromolecules are useful for diagnostics and other research purposes, for modulating protein in organisms, and for the diagnosis, detection and treatment of other conditions susceptible to antisense therapeutics.

, ,

19 Claims, 2 Drawing Sheets